

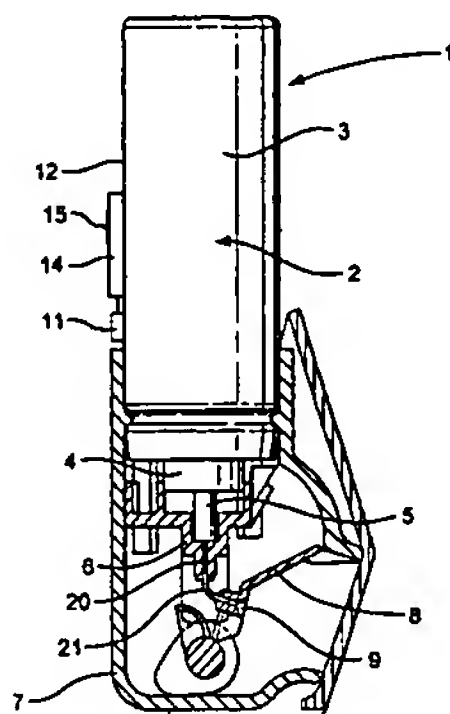
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- (71) Applicant and  
(72) Inventor: BACON, Raymond, John [GB/GB]; The Downer House, Aldworth, Emsworth, Hampshire PO10 8QT (GB).

[Continued on next page]

(54) Title: DISPENSER FOR MEDICAMENT



(57) Abstract: A dispenser (1) has a source of medicament (2) with a can (3) and a metered dose valve (4). The outlet stem (5) of the valve is fitted to a junction block (6) in a body (7) of the dispenser to which a breath actuated dose release mechanism (8) is connected for release of the medicament through a nozzle (9). The can has an acoustic transducer (11) attached to its side wall (12), in intimate acoustic contact. An electronic package (14), comprising a detection circuit and a driver for a LCD display (15) are also mounted on the can wall, the transducer being connected to the circuit. The detection circuit is arranged to decrement a count shown on the LCD by one for every detection of noise of dispensing/metering of a dose from the can. The detection circuit is programmed to compare the spectrum of the noise with a known dose dispensing spectrum and to decrement the counter according to the comparison. The counter starts at the manufacturer's recommended number of doses from the can, whereby the user can see how many doses are left to be dispensed and when a new dispenser is about to be needed.

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## DISPENSER FOR MEDICAMENT

The present invention relates to a dispenser for a medicament or the like to be inhaled as successive doses having a counter for such doses and to a counter therefor.

By way of illustration, in my prior International Patent Application, PCT/GB98/00770, at least as amended on entry into the European Regional Phase, there is described and claimed:

A dispenser for a gaseous, gas borne or droplet substance, the dispenser including:

- a body having a mouthpiece with an inhalation/insufflation orifice at its end;
- a junction in the body for a source of gas or evaporable liquid comprising or containing the said substance (the source being carried by the body); and
- a breath actuable valve, for controlling the release of said gas or liquid, comprising:
  - a valve inlet connected to the junction;
  - a valve outlet;
  - a flexible tube extending from the junction, between the inlet and the outlet, for receiving the said gas or liquid, the tube having a portion which is movable between a closed position in which the tube is kinked for closure of the valve and an open position in which the tube is un-kinked for opening of the valve; and
  - a movable member, for moving the movable portion of the tube to control its kinking, and being movably mounted in the body for movement by the act of inhalation from a rest position towards the orifice – or at least in the direction of air flow through the dispenser;
- the tube being kinked to an obturating extent when the movable member is in a rest position and un-kinked when the movable member is moved on inhalation for release of the gas or liquid.

The source of such a dispenser will typically be a pressurised can, having a valve for metering the dose of substance to be dispensed on each inhalation; although it is conceivable that this or a similar dispenser will dispense a dose which is not

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metered by the source, but rather is metered by the dispenser by filling of the duct between the source and the dispenser's valve on release of substance from the source.

There has been increasing interest in counting the number of doses dispensed from the source, for instance to alert the user to the number of doses which have been dispensed or more usually of the number of doses of known composition or strength which can still be dispensed.

The object of the present invention is to provide an improved counter for such a source and dispenser.

According to a first aspect of the invention, there is provided c, the dispenser comprising:

- a dispenser body having an inhalation passage leading to a mouthpiece;
- a pressurised source of the medicament arranged at an up-stream end of the inhalation passage;
- a junction in the body for receiving a stem of the pressurised source and directing the pressurised medicament for inhalation;
- means for controlling release of doses from the pressurised source;
- a transducer for detecting gaseous flow within the dispenser associated with release of a dose for inhalation; and
- a counter arranged to be incremented or decremented in accordance with each flow detection by the transducer.

In one embodiment, the source is a metered dose can and the junction includes a simple nozzle for directing the released dose towards the mouthpiece, the dose being released on depression of the can in the body. The transducer is an acoustic transducer arranged to detect noise of gas flow from the can on release of a dose.

Equally it is envisaged that an acoustic transducer detecting gas flow from the can could be provided in a breath actuated dispenser for detecting the preliminary release of the dose from the can prior to breath actuated release.

Conveniently the counter is adapted to recognise the spectrum of a particular gas flow in use of the dispenser. The acoustic transducer can be positioned in acoustic contact with the can, preferably at its side wall, for detecting gas flow to or from a metering chamber in the can. Alternatively, the acoustic transducer can be positioned in acoustic contact with the body of the dispenser for detecting gas flow to or from an intermediate chamber in dispenser.

In another embodiment, the source is a metered dose can, the junction provides connection to a breath actuated dose release mechanism. The transducer is a detector arranged to detect gas flow on inhalation causing the mechanism to release the dose following preliminary release of the dose from the can.

In accordance with a particular feature of the invention, the counter is associated with an indicator, typically audio or visual, for indicating a period of time from dose release during which the user should continue to inhale for drawing the medicament into his lungs or at least hold his breath with the medicament in his lungs to allow it to settle onto the lung lining.

In accordance with another feature, the counter is removably mounted on the dispenser, whereby it can be fitted to another dispenser after the can of a first dispenser has released its prescribed number of doses.

Again, the detector may be arranged to change the state of the counter from quiescent to active on detection of a first event and increment/decrement the counter on detection of a second event. The detector can comprise a single transducer for such two stage action or two transducers can be provided – one for each stage.

According to another aspect of the invention there is provided a counter and transducer for a dispenser according to the first aspect of the invention, the counter and transducer comprising:

- a transducer for detecting gaseous flow within the dispenser associated with release of a dose for inhalation; and

The detection circuit is arranged to decrement a count shown on the LCD by one for every detection of noise of dispensing/metering of a dose from the can. In order to avoid a spurious noise being detected as an indication of a dose, the detection circuit is programmed to compare the spectrum of the noise with a known dose dispensing spectrum and to decrement the counter according to the comparison. The counter starts at the manufacturer's recommended number of doses from the can, whereby the user can see how many doses are left to be dispensed and when a new dispenser is about to be needed.

It should be noted that the breath actuated dose release mechanism of the dispenser of Figure 1 and 2 plays no part in the operation of the dispenser. Rather, the transducer is arranged to detect the noise of a dose being dispensed from the can's internal metering chamber (not shown in detail) to the release mechanism 8, the metering chamber being a standard feature of such a can having a metered dose valve 4 and an outlet stem 5. The arrangement being such that on depression of the stem, a dose held in the metering chamber is released through the stem, and on release of the stem, it is resiliently returned and the metering chamber is replenished. For this the can needs to be inverted as it is in use. This is of particular note in respect of the term depression of the stem, which in this context involves an upwards movement of the stem inwards of the can.

It should also be noted that the transducer – or rather the acoustic recognition firmware in the counter – could be arranged to recognise the noise of the metering chamber refilling. However, it is believed that the noise of it emptying is more distinctive and more readily recognised. It is also conceivable that the firmware should recognise both noises and decrement the counter only on occurrence of every other recognised noise.

Further variants can be envisaged for this embodiment. For instance, the counter can be arranged to increment the displayed count towards a recommended maximum. Also, the transducer, circuitry and display can be arranged elsewhere, such as on the body 7 for detection of flow into an intermediate chamber 20 formed between the can stem 5 and a kink valve 21 in the breath actuated mechanism,

- a counter arranged to be incremented or decremented in accordance with each flow detection by the transducer.

The counter can be arranged merely to count. However, it is envisaged that other functions can also be performed, particularly initiation of a timer to time a delay between inhalation and breath release, to provide time for drug deposition in the lungs.

To help understanding of the invention, a specific embodiment thereof will now be described by way of example and with reference to the accompanying drawings, in which:

Figure 1 is a partially cut-away side view of a dispenser having a counter of the invention;

Figure 2 is an end view of the dispenser;

Figure 3 (a) and Figure 3 (b) are perspective views of a second dispenser according to the invention with its cover respectively closed and open;

Figure 4 (a) to Figure 4 (d) are cross-sectional views of the second dispenser respectively in closed, half-cock, cocked and dispenser states.

Referring to Figures 1 & 2, the dispenser 1 has a source of medicament 2 with a can 3 and a metered dose valve 4. The outlet stem 5 of the valve is fitted to a junction block 6 in a body 7 of the dispenser to which a breath actuated dose release mechanism 8 is connected for release of the medicament through a nozzle 9. Since the mechanism forms no part of the present invention, it will not be described in detail here. It could be configured as in my International Patent Application, PCT/GB01/03313, indeed Figures 1 and 2 are based on the second embodiment (Figure 6 et seq.) of that disclosure.

The can has an acoustic transducer 11 attached to its side wall 12, in intimate acoustic contact. An electronic package 14, comprising a detection circuit and a driver for a LCD display 15 are also mounted on the can wall, the transducer being connected to the circuit. It is believed that the selection of the transducer, detection circuit, driver and display will all be within the capabilities of the man skilled in the art, and as such will not be described here in detail.

whereby the transducer on the body can recognise the noise of the pressurised dose being dispensed through the nozzle.

Referring to Figures 3(a) to 4(d), a further metered dose inhaler is thereshown, this is based on the third embodiment of my International Patent Application, PCT/GB01/03313, namely that of Figures 13 to 17 thereof. The dispenser thereshown has a body 201 with a mouthpiece 202 and a pivoted mouthpiece cover 203. The mouthpiece is formed as an aperture 2021 in a separate body part 2012 clipped to a main body part 2011. The mouthpiece part is cutaway 2014 with respect to the medicament can 211 fitted to the body, to define an air inlet exposed by the cover when this is open. The cover is pivoted about an axis 204 low in the body at the joint between the two body parts.

The can 211 is held in an opening 210 at the upper end of the main body part 2011 by a pair of location pins 2101 in openings 2102 in sidewalls of the body part. The pins positively locate the can with respect to the body by engagement in a groove 215 formed in the can for retaining its closure collar. Thus the can and the body 201 are rigidly connected. Also moulded inside the body are internal ribs 216. A junction member 217 is slidably accommodated in the body with the ribs engaging in grooves 218 in its periphery. Centrally, the junction member has a socket 220 for an outlet stem 221 of the can. The socket is continued by a passage 222, which has a thin wall, kinkable portion 223 and a nozzle end 224. This is in a movable outlet member 225 of a valve part 2172 of the junction member. The main part 2171 of the junction member 217 and the valve part 2172 are connected by a living hinge. To both sides of the outlet member are provided flats 2251, which form the base for a breath actuation flap 228 and are connected to the living hinge portions at the outside ends of the flats. A lug 227 depends from the outlet member for engagement with a cam finger 232 extending from between a pair of cam lobes 2052.

To either side of the socket 220 of the junction member, it has two depending fingers 230 arranged to co-operate with the cam lobes at their distal ends 2301. These abut – under the force of the internal spring (not shown) of the can, and with the interposition of two leaves 2019 – the cam lobes 2052.

The action of the dispenser is as follows:

On opening of the cover, by swinging about the axis of the cam member, the cam finger 232 acts on the lug 227 to move the outlet member to a position where the flap 228 is lifted and the kinkable portion 223 is closed. In this position, the kink tube is kinked and will not pass a dose about to be released into it.

Further opening of the cover causes the cam lobes 2051, 2052 to lift the junction member and stem 221 towards the can. This releases a dose from the can into the kinked valve, which retains it. The cover stops by abutment with body and the mechanism is cocked and primed for use.

Breathing in through the mouthpiece draws air across the flap, that is round its edges from the air inlet 2014, with a pressure differential developing. Over-centre spring retention of the flap is overcome and it is sprung down to dispense the dose as the outlet nozzle points into the mouthpiece. This, it is free to do since the lug 227 is free of the cam finger 232 at this position.

The mechanism is reset by closure of the cover over the mouthpiece. The junction member drops under the action of the can valve spring and control of the main cam lobes. The cam finger 232 and the flap lug 227 engage on their rear faces, that is the faces opposite from the those which cause lifting of the flap on opening of the cover. Such engagement is unwanted and the faces are provided with complementary wedge shapes – as shown in Figure 12 – whereby the lobe and lug deflect side ways and pass each other. This deflection causes a drag on the lug and keeps the flap in its open position. To ensure that the lobe and lug re-engage for next use, each being thin for deflection, their front faces are provided with complementary V edge.

The action of closing the cover causes the cam fingers (not shown) to engage the flap fingers – should the flap be pivoted towards the can – and pivot the flap to its position in which the nozzle 224 is directed towards the (closed) mouthpiece and the kink tube is unkinked.

Should the inhaler not be used, the act of closure releases the flap, and rests the breath actuated mechanism. This releases the dose against the inside of the cover. It is a dose that is lost. The counter counts this since the deflection of the second flap to its quiescent state 1063 causes a signal to be passed to the counter. The latter decrements its displayed count.

In a non-illustrated variant, the piezo-electric strip 106 is arranged not to be deflected, but to experience a change in temperature on inhalation and thus decrement the counter.

In an other non-illustrated variant, the transducers are proximity detectors arranged to detect movement of the cover and the breath actuated flap.

A counter 101 is removably mounted on the can 211. It comprises a ring 102 sized to grip the can, yet removable in case of need to refit it to another can. It has an LCD display 103, associated with a counting and timing circuit mounted in the ring. The ring carries two transducing elements. The first is an arched piece of piezo-electric film 105 abutted by the edge of the cover and held flat when the cover is closed. The second is a second piezoelectric film 106 extending through the air inlet 2014, to be abutted by the breath actuation flap 228 when the flap is set to the cocked position in Figure 4(c). This abutment flexes the film from a flat state to which it has a tendency to return.

On opening of the cover, the first film 105 is free to resume its arched state 1051 and in doing so sends a “wake-up” signal to the counter. The flap 228 is set to its cocked state, deflecting the second film 106 to a curved state 1061. This causes it to send a signal to the counter. This is ignored or can be used as an additional wake-up signal or indeed in the possible absence of the film 105, this signal can be the wake-up signal for the counter. On inhalation, two effects combine to flex the second film 106. Firstly the deflection imposed by the flap is removed. Second inhalation air flexes the film to an oppositely curved state 1062. This flexure causes it to generate a “count” signal causes the counter to decrement by one the count of doses shown on the LCD display 103.

Further, the count signal causes the counter to illuminate an LED 108, indicating to the user to keep inhaling or at least to not exhale. The LED can be illuminated with a red colour. At the end of an inhalation period, say two seconds, sufficient for the medicament to reach the user's lungs and begin to settle out of suspension, the LED is caused to change colour to green, indicating to the user that exhalation is now allowed. The LED can be supplemented or replaced by a two tone buzzer 109 or by flashing or other indication from the LCD display.

After use, the cover is closed, flattening the first film 105. The second film 106 takes up a quiescent flat state 1063. It should be noted that second film detects both cocking of the dispenser, when deflected from the quiescent state 1063 to the deflected state 1061 imposed by the breath actuation flap 1062, and inhalation gas flow when deflected as a vane by this flow to the deflected state 1062.

#### CLAIMS:

1. A dispenser for a medicament or the like to be inhaled as successive doses, the dispenser comprising:

- a dispenser body having an inhalation passage leading to a mouthpiece;
- a pressurised source of the medicament arranged at an up-stream end of the inhalation passage;
- a junction in the body for receiving a stem of the pressurised source and directing the pressurised medicament for inhalation;
- means for controlling release of doses from the pressurised source;
- a transducer for detecting gaseous flow within the dispenser associated with release of a dose for inhalation; and
- a counter arranged to be incremented or decremented in accordance with each flow detection by the transducer.

2. A dispenser as claimed in claim 1, wherein:

- the source is a metered dose can having a metering chamber;
- the dose release controlling means is a valve in the can arranged to release a dose from the metering chamber on depression of the stem towards the can;
- the junction includes a simple nozzle for directing the released dose towards the mouthpiece, the dose being released on depression of the can in the body; and
- the transducer is an acoustic transducer arranged to detect noise of gas flow within or from the can on release of a dose.

3. A dispenser as claimed in claim 1, wherein:

- the source is a metered dose can having a metering chamber;
- the dose release controlling means is a breath actuated release mechanism downstream of the junction, the dose being released from the mechanism on inhalation;
- the junction includes a connection from the can to the breath actuated release mechanism, the dose being released on depression of the can in the body; and
- the transducer is an acoustic transducer arranged to detect noise of gas flow within or from the can on release of a dose.



4. A dispenser as claimed in claim 2 or claim 3, wherein the counter is adapted to recognise the spectrum of a particular gas flow in use of the dispenser and to increment/decrement the counter in response to such recognition.
5. A dispenser as claimed in claim 2, claim 3 or claim 4, wherein the acoustic transducer is positioned in acoustic contact with the can, preferably at its side wall, for detecting gas flow to or from a metering chamber in the can.
6. A dispenser as claimed in claim 2, claim 3 or claim 4, wherein the acoustic transducer is positioned in acoustic contact with the body of the dispenser for detecting gas flow in a gas passage down-stream of the can.
7. A dispenser as claimed in claim 1, wherein:
- the source is a metered dose can having a metering chamber;
  - the dose release controlling means is a breath actuated release mechanism downstream of the junction, the dose being released from the mechanism on inhalation;
  - the junction includes a connection from the can to the breath actuated release mechanism, the dose being released on depression of the can in the body; and
  - the transducer is arranged to detect inhalation gas flow within the body.
8. A dispenser as claimed in claim 7, wherein:
- the breath actuated release mechanism includes a release flap acted on by inhalation flow and
  - the transducer is arranged to detect movement of the flap.
9. A dispenser as claimed in claim 7, wherein the transducer is a flow transducer arranged to detect flow of inhalation gas past it.
10. A dispenser as claimed in claim 9, wherein the transducer comprises a vane extending into the inhalation passage and adapted to detect a flow induced pressure differential across the vane.
11. A dispenser as claimed in claim 10, wherein the vane is of piezoelectric film material.
12. A dispenser as claimed in claim 7, wherein the transducer is a temperature transducer arranged to detect flow of gas past it as a change in temperature of gas in contact with it.
13. A dispenser as claimed in claim 12, wherein the temperature transducer is of piezoelectric film material.

14. A dispenser as claimed in any preceding claim, wherein the counter is associated with an indicator, preferably audio or visual, for indicating a period of time from dose release during which the user should continue to inhale or at least hold his breath for drawing the medicament into his lungs.
15. A dispenser as claimed in any preceding claim, wherein the counter is removably mounted on the dispenser, whereby it can be fitted to another dispenser after the can of a first dispenser has released its prescribed number of doses.
16. A dispenser as claimed in any preceding claim, wherein the counter is arranged to change the state of the counter from quiescent to active on detection of a first event and increment/decrement the counter on detection of a second event.
17. A dispenser as claimed in claim 16, comprising a single transducer for detecting such first events and second events.
18. A dispenser as claimed in claim 16, comprising two transducers, one for detecting such first events and the other for detecting such second events.
19. A dispenser as claimed in claim 18, including a mouthpiece cover and wherein the one transducer is adapted and arranged to detect opening of the cover and the other transducer is the gas flow detection transducer.
20. A dispenser as claimed in claim 19 as appendant to claim 3 or claim 7 or any other claim appendant thereto, wherein:
- the cover is adapted and arranged to release a dose from the source to the breath actuated release mechanism on opening of the cover and to release the dose therefrom on closure of the cover without breath actuated dose release and
  - the counter is arranged to increment/decrement in the event of such closure.
21. A counter and transducer for a dispenser as claimed in any preceding claim, the counter and transducer comprising:
- a transducer for detecting gaseous flow within the dispenser associated with release of a dose for inhalation; and
  - a counter arranged to be incremented or decremented in accordance with each flow detection by the transducer.

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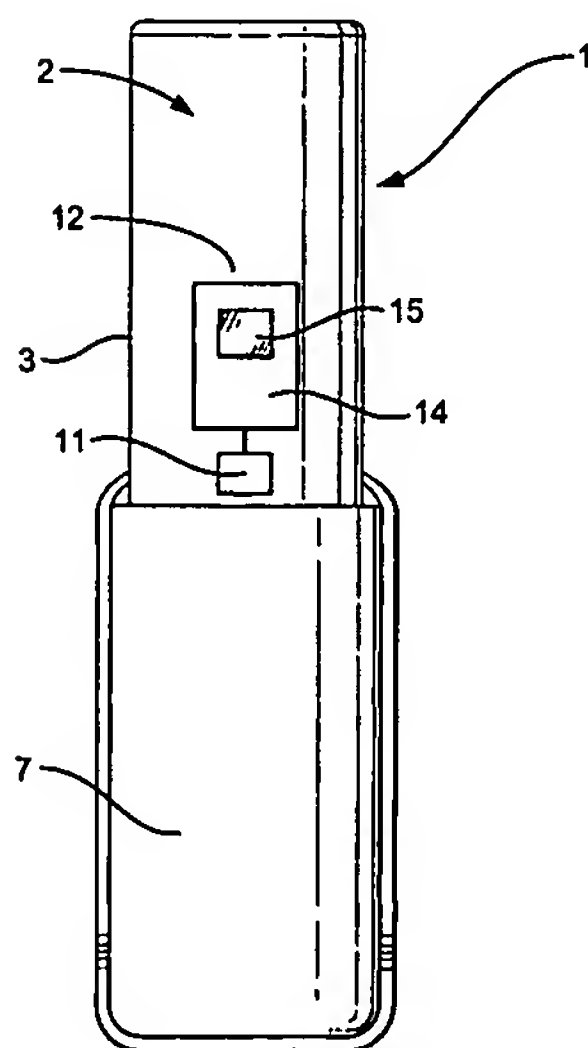


Fig. 1

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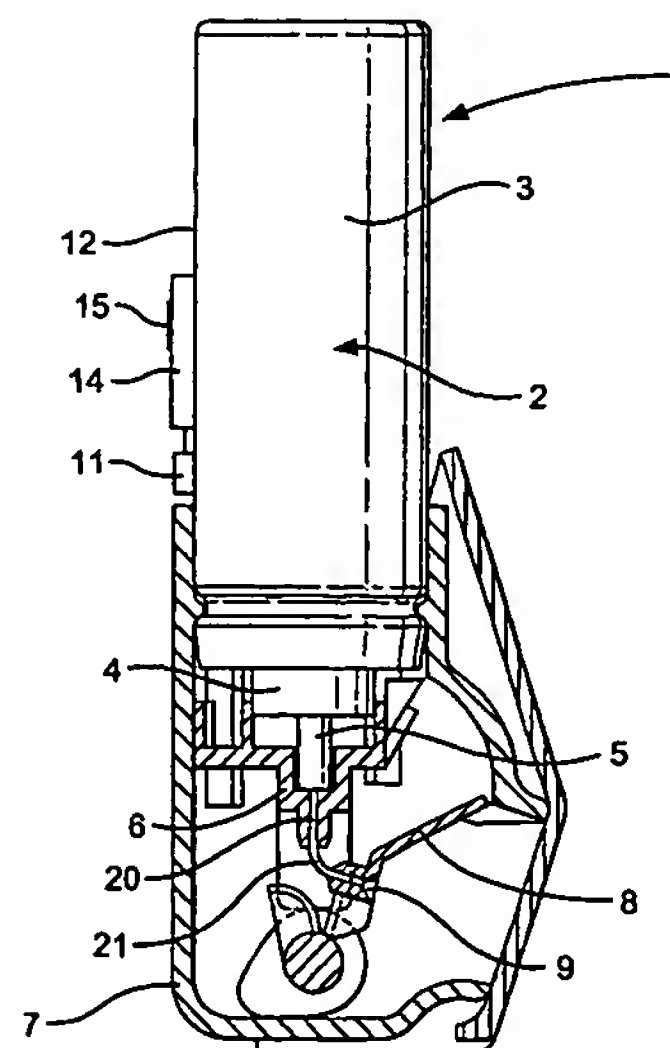
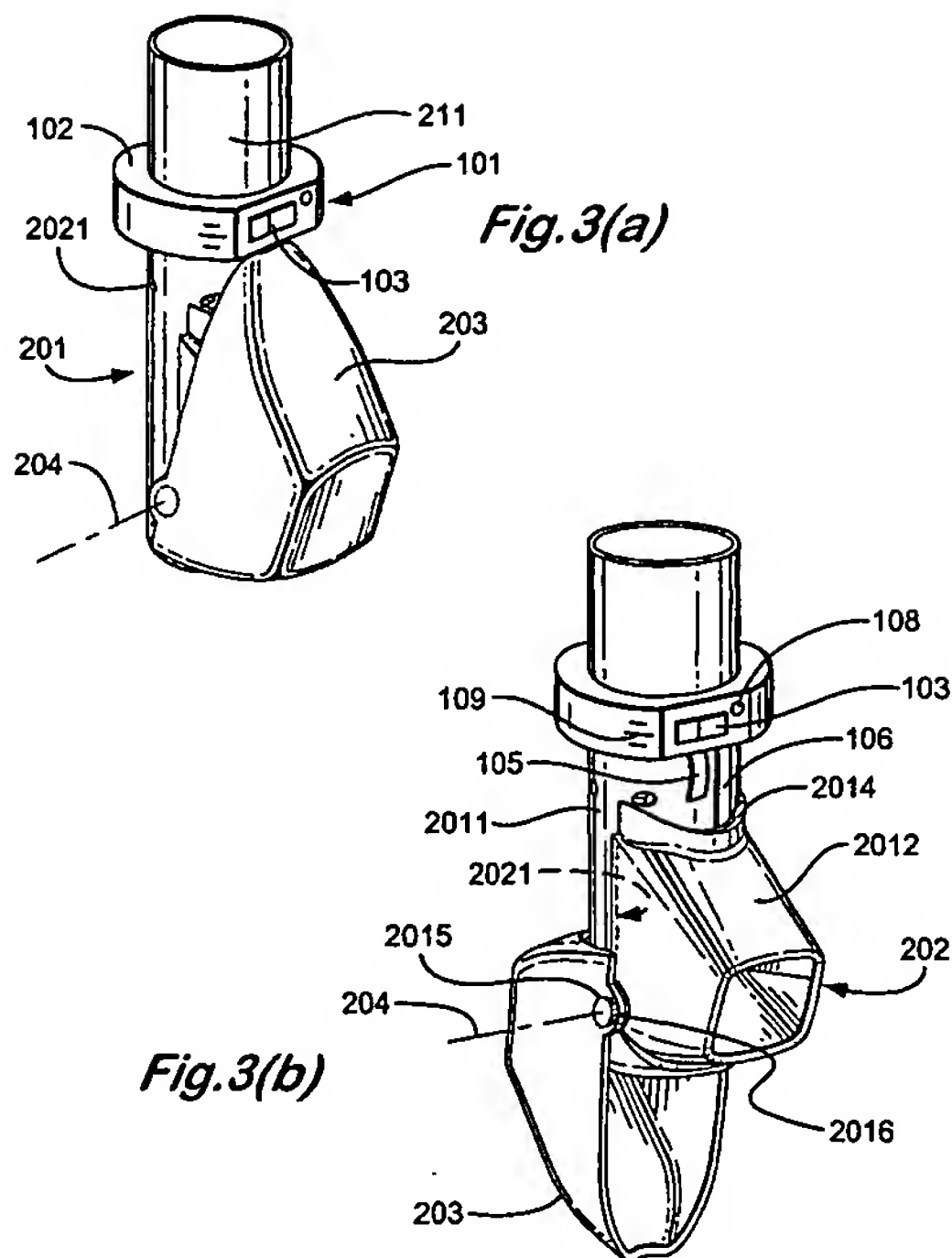


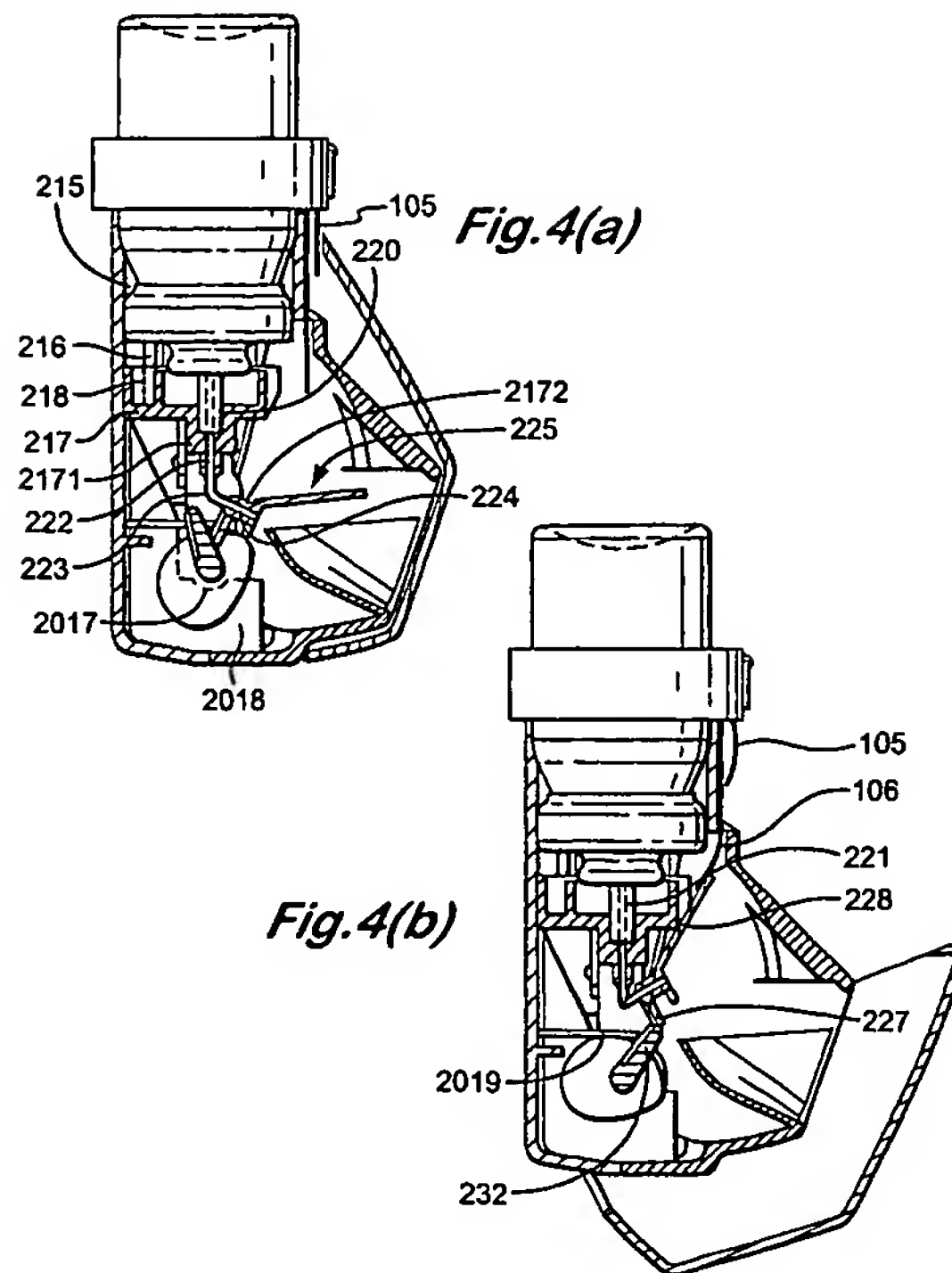
Fig. 2

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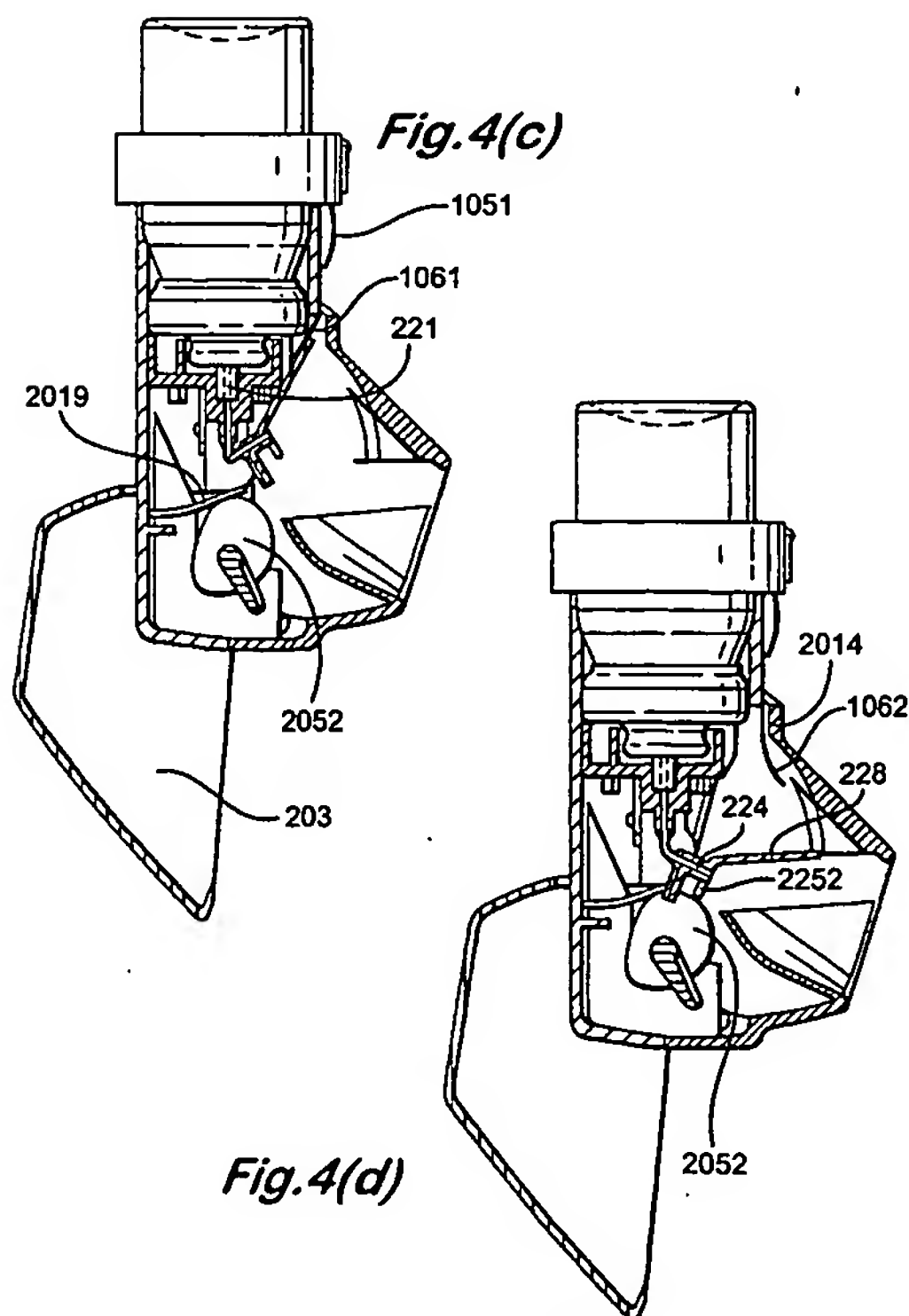
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## INTERNATIONAL SEARCH REPORT

PCT/GB 02/00285

A. CLASSIFICATION OF SUBJECT MATTER  
IPC 7 A61M15/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 A61M 610K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 794 612 A (WACHTER ALLAN ET AL) 18 August 1998 (1998-08-18) the whole document	1, 2, 4-6, 14-19, 21 3, 7-13, 20
A	US 5 544 647 A (EBELING FREDERICK A ET AL) 13 August 1996 (1996-08-13) abstract; figures column 1, line 6-12 column 2, line 52 - column 3, line 5 column 7, line 41 - column 8, line 26	1-19  20
Y		
A		

☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

24 April 2002

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Name and mailing address of the ISA

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 85504 Munich, Germany  
 Tel: (+31-70) 940-2000, Tx: 31 851 spe nl  
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INTERNATIONAL SEARCH REPORT		PCT/GB 02/00285
C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 99 64095 A (MICRODOSE TECHNOLOGIES INC) 16 December 1999 (1999-12-16) abstract	21
Y		1,2,4-6, 14-19
A	page 8, line 4 -page 9, line 8 page 13, line 28 -page 14, line 6 page 19, line 26 -page 20, line 7 page 24, line 7-18 page 24, line 29 -page 25, line 12	3,7-13, 20
Y	US 5 415 161 A (RYDER STEVEN L) 16 May 1995 (1995-05-16) abstract; figures column 1, line 6-9 column 2, line 17-24 column 2, line 50-68 column 4, line 19-28 column 4, line 48-68 column 5, line 37-59	3,7-13
X	US 6 152 130 A (FLEMING SCOTT ET AL) 28 November 2000 (2000-11-28) abstract; figures column 1, line 6-11 column 2, line 16-47	21
A	column 3, line 11 -column 4, line 2 column 4, line 63 -column 5, line 10 column 5, line 31 -column 6, line 7	1-20
A	US 5 839 429 A (MARNFELDT NILS GOERAN ET AL) 24 November 1998 (1998-11-24) abstract; figures column 1, line 7-13 column 1, line 38-47 column 2, line 9-21 column 2, line 30-41 column 2, line 58-67	1-21
A	US 4 664 107 A (WASS ANTHONY C L) 12 May 1987 (1987-05-12) abstract; figures	3,7-13

Form PCT/ISA/210 (continuation of second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT				PCT/GB 02/00285	
Patent document cited in search report		Publication date	Patent family member(s)		Publication date
US 5794612	A	18-08-1998	US	6085742 A	11-07-2000
US 5544647	A	13-08-1996	AT	211399 T	15-01-2002
			AU	687290 B2	19-02-1998
			AU	4236096 A	19-06-1996
			CA	2181789 A1	06-06-1996
			DE	69524865 D1	07-02-2002
			EP	0746366 A1	11-12-1996
			JP	9508845 T	09-09-1997
			NO	963134 A	28-08-1996
			NZ	297246 A	22-09-1997
			WO	9616686 A1	06-06-1996
			US	5622163 A	22-04-1997
			ZA	9510129 A	06-06-1996
WO 9964095	A	16-12-1999	US	5960609 A	05-10-1999
			US	6142146 A	07-11-2000
			US	6152130 A	28-11-2000
			AU	4683199 A	30-12-1999
			BR	9911098 A	29-01-2002
			CN	1320054 T	31-10-2001
			EP	1083952 A2	21-03-2001
			WO	9964095 A2	16-12-1999
US 5415161	A	16-05-1995	NONE		
US 6152130	A	28-11-2000	AU	4683199 A	30-12-1999
			BR	9911098 A	29-01-2002
			CN	1320054 T	31-10-2001
			EP	1083952 A2	21-03-2001
			WO	9964095 A2	16-12-1999
US 5839429	A	24-11-1998	AU	2154595 A	17-10-1995
			WO	9526212 A1	05-10-1995
US 4664107	A	12-05-1987	AU	578915 B2	10-11-1988
			AU	3558384 A	22-05-1985
			CA	1236359 A1	10-05-1988
			DE	3470844 D1	09-06-1988
			EP	0147028 A1	03-07-1985
			FI	852490 A ,B,	24-06-1985
			JP	5048138 B	20-07-1993
			JP	61500254 T	20-02-1986
			WO	8501880 A1	09-05-1985

Form PCT/ISA/210 (second family annex) (July 1992)